Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

- Claims 1-21 (Canceled)
- Claim 22 (Previously Presented) A purified polypeptide capable of forming antigen binding structures with specificity for Rhesus D antigens comprising a V_H region having SEQ ID NO: 58 or a functional variant thereof and a V_L region, wherein said functional variant of the V_H region is SEQ ID NO: 2, SEQ ID NO: 6, SEQ ID NO: 10, SEQ ID NO: 14, SEQ ID NO: 18, SEQ ID NO: 22, SEQ ID NO: 26, SEQ ID NO: 30, SEQ ID NO: 34, SEQ ID NO: 38, SEQ ID NO: 42, SEQ ID NO: 46, SEQ ID NO: 50, SEQ ID NO: 54, or SEQ ID NO: 62.
- Claim 23. (Previously Presented) The polypeptide of claim 22, wherein the V_L region is SEQ ID NO: 4, SEQ ID NO: 8, SEQ ID NO: 12, SEQ ID NO: 16, SEQ ID NO: 20, SEQ ID NO: 24, SEQ ID NO: 28, SEQ ID NO: 32, SEQ ID NO: 36, SEQ ID NO: 40, SEQ ID NO: 44, SEQ ID NO: 48, SEQ ID NO: 52, SEQ ID NO: 56, SEQ ID NO: 60, or SEQ ID NO: 64.
- Claim 24. (Canceled)
- Claim 25. (Previously Presented) The polypeptide of claim 22, wherein the polypeptide is an antigen binding Fab fragment.
- Claim 26. (Previously Presented) The polypeptide of claim 22, wherein the polypeptide is an immunoglobulin specific for a Rhesus D antigen.
- Claim 27. (Presently Presented) The polypeptide of claim 26, wherein the immunoglobulin comprises at least one defined isotype selected from the group consisting of IgG1, IgG2, IgG3, and IgG4.
- Claim 28. (Previously Presented) A recombinant polynucleotide which encodes the polypeptide of claim 22.

- Claim 29. (Previously Presented) A pharmaceutical composition comprising at least one polypeptide of claim 22.
- Claim 30. (Previously Presented) A pharmaceutical composition comprising at least one immunoglobulin of claim 26.
- Claim 31. (Previously Presented) A diagnostic composition for Rhesus D typing comprising at least one polypeptide of claim 22.
- Claim 32. (Previously Presented) A diagnostic composition for Rhesus D typing comprising at least one immunoglobulin of claim 26.
- Claim 33. (Currently Amended) A purified polypeptide capable of forming antigen binding structures with specificity for Rhesus D antigens comprising a V_H region and a V_L region having SEQ ID NO: 60 or a functional variant thereof, wherein said functional variant of the V_L region is SEQ ID NO: 4, SEQ ID NO: 8, SEQ ID NO: 12, SEQ ID NO: 16, SEQ ID NO: 20, SEQ ID NO: 24, SEQ ID NO: 28, SEQ ID NO: 32, SEQ ID NO: 36, SEQ ID NO: 40, SEQ ID NO: 44, SEQ ID NO: 48, SEQ ID NO: 52, SEQ ID NO: 56, or SEQ ID NO: 64.
- Claim 34. (Previously Presented) The polypeptide of claim 33, wherein the V_H region is SEQ ID NO: 2, SEQ ID NO: 6, SEQ ID NO: 10, SEQ ID NO: 14, SEQ ID NO: 18, SEQ ID NO: 22, SEQ ID NO: 26, SEQ ID NO: 30, SEQ ID NO: 34, SEQ ID NO: 38, SEQ ID NO: 42, SEQ ID NO: 46, SEQ ID NO: 50, SEQ ID NO: 54, SEQ ID NO: 58, or SEQ ID NO: 62.
- Claim 35. (Canceled)
- Claim 36. (Previously Presented) The polypeptide of claim 33, wherein the polypeptide is an antigen binding Fab fragment.
- Claim 37. (Previously Presented) The polypeptide of claim 33, wherein the polypeptide is an immunoglobulin specific for a Rhesus D antigen.

- Claim 38. (Previously Presented) The polypeptide of claim 37, wherein the immunoglobulin comprises at least one defined isotype selected from the group consisting of IgG1, IgG2, IgG3, and IgG4.
- Claim 39. (Previously Presented) A recombinant polynucleotide which encodes the polypeptide of claim 33.
- Claim 40. (Previously Presented) A pharmaceutical composition comprising at least one polypeptide of claim 33.
- Claim 41. (Previously Presented) A pharmaceutical composition comprising at least one immunoglobulin of claim 37.
- Claim 42. (Previously Presented) A diagnostic composition for Rhesus D typing comprising at least one polypeptide of claim 33.
- Claim 43. (Previously Presented) A diagnostic composition for Rhesus D typing comprising at least one immunoglobulin of claim 37.
- Claim 44. (Withdrawn) A method for preventing or treating a hematologic disorder in a subject comprising administering to the subject a pharmaceutical composition comprising at least one of the following
 - (a) a purified polypeptide capable of forming antigen binding structures with specificity for Rhesus D antigens comprising a V_H region having SEQ ID NO: 58 or a functional variant thereof and a V_L region;
 - (b) a purified polypeptide capable of forming antigen binding structures with specificity for Rhesus D antigens comprising a V_H region and a V_L region having SEQ ID NO: 60 or a functional variant thereof;
 - (c) an immunoglobulin specific for a Rhesus D antigen comprising a purified polypeptide having a V_H region having SEQ ID NO: 58 or a functional variant thereof and a V_L region;

(d) an immunoglobulin specific for a Rhesus D antigen comprising a purified polypeptide having a V_H region and a V_L region having SEQ ID NO: 60 or a functional variant thereof;
wherein the hematologic disorder is haemolytic disease of the newborn (HDN), immune thrombocytopenic purpura (ITP), or mistransfusion of Rhesus incompatible blood.